



To the European Patent Office
Entry into the European phase (EPO as designated or elected Office)

European application number	EP04804263.4
PCT application number	PCT/EP2004/014669
PCT publication number	WO2005064871
Applicant's or representative's reference	P06481EP01
1. Applicant	
Particulars of the applicant(s) are contained in the international publication or were recorded by the International Bureau subsequent to the international publication.	
<input checked="" type="checkbox"/> Changes which have not yet been recorded by the International Bureau are set out here:	
Address for correspondence	
2. Representative 1	
This is the representative who will be listed in the Register of European Patents and to whom notifications will be made	
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Any additional representative(s) is/are listed here:	<input checked="" type="checkbox"/>
3. General Authorisation: An individual authorisation is attached. <input type="checkbox"/> A general authorisation has been registered under No: <input type="checkbox"/> A general authorisation has been filed, but not yet registered. <input type="checkbox"/> The authorisation filed with the EPO as PCT receiving Office expressly includes the European phase.	
4. Request for examination Examination of the application under Art. 94 EPC is hereby requested. The examination fee is being (has been, will be) paid. <input checked="" type="checkbox"/> Request for examination in an admissible non-EPO language: <input checked="" type="checkbox"/>	
5. Copies One or more additional sets of copies of the documents cited in the supplementary European search report are hereby requested. <input type="checkbox"/> Number of additional sets of copies	
6. Documents intended for proceedings before the EPO 6.1 Proceedings before the EPO as designated Office (PCT I) are to be based on the following documents:	

<p>the application documents published by the International Bureau (with all claims, description and drawings), where applicable with amended claims under Art. 19 PCT unless replaced by the amendments attached.</p> <p><i>Where necessary, clarifications should be attached as 'Other Documents'</i></p> <p>6.2 Proceedings before the EPO as elected Office (PCT II) are to be based on the following documents:</p> <p>the documents on which the international preliminary examination report is based, including any annexes unless replaced by the amendments attached.</p> <p><i>Where necessary, clarifications should be attached as 'Other Documents'</i></p> <p>If the EPO as International Preliminary Examining Authority has been supplied with test reports, these may be used as the basis of proceedings before the EPO.</p>	<input type="checkbox"/> <input type="checkbox"/> <input checked="" type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
<p>7. Translations</p> <p>Translations in one of the official languages of the EPO (English, French, German) are attached as crossed below:</p> <p>* <i>In proceedings before the EPO as designated or elected Office (PCT I + II):</i></p> <p>Translation of the international application (description, claims, any text in the drawings) as originally filed, of the abstract as published and of any indication under Rule 13bis.3 and 13bis.4 PCT regarding biological material</p> <p>Translation of the priority application(s)</p> <p>It is hereby declared that the international application as originally filed is a complete translation of the previous application (Rule 38(5) EPC)</p> <p>* <i>In addition, in proceedings before the EPO as designated Office (PCT I):</i></p> <p>Translation of amended claims and any statement under Art. 19 PCT, if the claims as amended are to form the basis for the proceedings before the EPO (see Section 6).</p> <p>* <i>In addition, in proceedings before the EPO as elected office (PCT II):</i></p> <p>Translation of annexes to the international preliminary examination report</p>	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
<p>8. Biological material</p> <p>The invention relates to and/or uses biological material deposited under Rule 28 EPC.</p> <p>The particulars referred to in Rule 28(1)(c) EPC (if not yet known, the depositary institution and the identification reference(s) [number, symbols, etc.] of the depositor) are given in the international publication or in the translation submitted under Section 7 on:</p> <p>page(s) / line(s)</p> <p>A copy of the receipt(s) of deposit issued by the depositary institution is attached</p> <p>will be filed at a later date</p> <p>A waiver of the right to an undertaking from the requester pursuant to Rule 28(3) EPC is attached.</p>	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
<p>9. Nucleotide and amino acid sequences</p> <p>The items required under Rules 5.2 and 13ter PCT and Rule 111(3) EPC have already been furnished to the EPO.</p> <p>The sequence listing as part of the description is attached in PDF format.</p> <p>The sequence listing does not include matter that goes beyond the content of the application as filed.</p> <p>In addition, the sequence listing data is attached in computer-readable form in accordance with WIPO Standard 25.</p> <p>The sequence listing data in computer-readable form in accordance with WIPO Standard 25 is identical to the sequence listing in PDF format.</p>	<input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/> <input type="checkbox"/>
<p>10. Designation fees</p> <p>10.1 It is currently intended to pay seven times the amount of the designation fee. The designation fees for all the EPC contracting states designated in the international application are thereby deemed to have been paid (Art. 2 No. 3 RFees).</p> <p>AT BE BG CH&LI CY CZ DE DK EE ES FI FR GB GR HU IE IS IT LT LU MC NL PL PT RO SE SI SK TR</p>	<input checked="" type="checkbox"/>

10.2 It is currently intended to pay fewer than seven designation fees for the following EPC contracting states designated in the international application:

10.3 It is requested that no communication under Rules 85a(1) or 69(1) need be notified in respect of the contracting states not indicated. If an automatic debit order has been issued, the EPO is authorised, on expiry of the basic period under Article 79(2), to debit seven times the amount of the designation fee. If less than seven states are indicated, the EPO shall debit designation fees only for those states, unless it is instructed to do otherwise before expiry of the basic period.

11. Extension of the European patent

This application is also considered as being a request for extension to all the non-contracting states to the EPC designated in the international application with which "extension agreements" were in force on the date of filing the international application. However, the extension only takes effect if the prescribed extension fee is paid.

It is currently intended to pay the extension fee for the following states:

12. List of enclosed documents

Description of document	Original file name	Assigned file name
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13. Debit from deposit account

Currency

EUR

The European Patent Office is hereby authorised, to debit from the deposit account with the EPO any fees and costs indicated on the fees page.

Deposit account number

28100039

Account holder

DR LUDWIG BRANN
PATENTBYRÅ AB

14. Reimbursements (if any) should be made to the following EPO deposit account:

Number and account holder

DR LUDWIG BRANN
PATENTBYRÅ AB, 28100039

15. Fees

	Total:	Factor applied	Fee schedule	Amount to be paid
			EUR	0.00

16. Annotations

16-1. Note (for EPO) (EP Phase)

Fees payable (Karolina Karlsson;
20.6.-6-21)

Owing to a technical problem in OLF
the correct fees will be paid by fax.

17. Signature(s) of applicant(s) or representative

Place:

Uppsala

Date:

22.June 2006

Signed by:

SE, Dr. Ludwig Brann Patentbyra AB, H.-A.
Svanfeldt 1948

Capacity:

(Representative)